

## WE CLAIM:

 (currently amended): A process for preparing a eephalosporin cefdinir of Formula II

Formula II

comprising the steps of:

reacting O-acetyl thioester of Formula I

Formula I

with a compound of Formula III in the presence of a base in suitable solvent

Formula III

wherein R' [[= H,]] represents H or any carboxyl protecting group or silyl group, converting to cefdinir by the removal of protecting group or groups.

- 2. (original): The process according to claim 1 wherein the said base can be organic base or an inorganic base.
- 3. (currently amended): The process according to claim 1 wherein the said organic base is an amine selected from the group consisting of triethylamine, N,N-diisopropylethylamine, tributylamine tri n-butylamine.
- 4. (original): The process according to claim1 wherein the said inorganic base is selected from the group consisting of sodium carbonate, sodium bicarbonate and mixtures thereof.
- 5. (original): The process according to claim 1 wherein the said solvent is selected from the group consisting of water, tetrahydrofuran, methylene dichloride and mixtures thereof.
- 6. (currently amended) The process according to claim 1 wherein the said reacting step is conducted at a temperature between 10°C and 25°C.
- 7. (currently amended) The process according to claim 1 wherein the said carboxyl protecting group is selected from the group consisting of p-methoxybenzyl, p-nitrobenzyl, and diphenylmethyl and trimethyl silyl.

(currently amended): The process according to claim 1 wherein the said to prepare
O-acetyl thioester of Formula I

is prepared by a process which comprises of

condensing (Z)-2-(2-amino-4-thiazolyl)-2-acetyloxyiminoacetic acid with bis(benzothiazol-2-yl)disulphide in the presence of triphenylphosphine and a base in a suitable solvent.

- 9. (original): The process according to claim 8 wherein the said base is selected from the group consisting of tributylamine, triethylamine and mixtures thereof.
- 10. (original): The process according to claim 8 wherein the said solvent is selected from the group consisting of methylene chloride, chloroform, tetrahydrofuran, acetonitrile and mixtures thereof.
- 11. (original): The process according to claim 8 wherein the said reacting step is conducted at a temperature between 0°C and 35°C.